## **Claims**

## What is claimed is:

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- 5 1. An isolated polypeptide having glucose isomerase activity, selected from the group consisting of:
  - (a) a polypeptide having an amino acid sequence which has at least 95% identity with amino acids for the mature polypeptide of SEQ ID NO:2;
  - (b) a variant of the polypeptide having an amino acid sequence of SEQ ID NO:2 comprising a substitution, deletion, and/or insertion of one or more amino acids;
    - (c) a fragment of (a) that has glucose isomerase activity; and
    - (d) a polypeptide having a pH optimum in the range of 5.7 to 6.3 at 60 °C, a pH optimum in the range of 6.1 to 6.7 at 90 °C and a temperature optimum of above 90 °C.
- 15 2. The polypeptide of claim 1, comprising the amino acid sequence of SEQ ID NO:2.
  - 3. The polypeptide of claim 1, which has a pH optimum in the range of 5.9 to 6.1 at 60 °C.
- 20 4. The polypeptide of claim 1, which has a pH optimum in the range of 6.3 to 6.5 at 90 °C.
  - 5. The polypeptide of claim 1, which has a pH optimum of about 6 at 60 °C, a pH optimum of about 6.4 at 90 °C and a temperature optimum of 90 to 100 °C.
  - 6. The polypeptide of claim 1, which has a specific activity on fructose substrate of at least 15 unitF/mg.
  - 7. The polypeptide of claim 1, which is encoded by the nucleic acid sequence contained in plasmid pBSK1 or plasmid pBSK2.

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- 8. The polypeptide of claim 1 being derived from the strain Streptomyces sp. SK (GI SK).
- 9. The strain Streptomyces sp. SK expressing genes encoding a polypeptide of claim 1.
- 10. A glucose isomerase GI in which a residue other than alanine at a position equivalent to position 100, 102 and/or 109 of GI SK has/have been replaced by alanine.
- 11. The glucose isomerase of claim 10, wherein the corresponding naturally occurring GI is derived from a microorganism of the order Actinomycetales.
  - 12. The glucose isomerase of claim 11, wherein the corresponding naturally occurring GI is derived from a microorganism of *Streptomyces*.
  - 13. An isolated nucleic acid sequence comprising a nucleic acid sequence which encodes the polypeptide of claim 1.
  - 14. An isolated nucleic acid sequence comprising a nucleic acid sequence having at least one mutation in the mature polypeptide coding sequence of SEQ ID NO:1, in which the mutant nucleic acid sequence encodes a polypeptide consisting of amino acids of SEQ ID NO:2.
- 15. A nucleic acid construct comprising the nucleic acid sequence of claim 13 operably linked to one or more control sequences that direct the production of the polypeptide in a suitable expression host.
  - 16. A recombinant expression vector comprising the nucleic acid construct of claim 15.
  - 17. A recombinant host cell comprising the nucleic acid construct of claim 15.
  - 18. A method for producing a mutant nucleic acid sequence, comprising: (a) introducing

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at least one mutation into the mature polypeptide coding sequence of SEQ ID NO:1; and (b) recovering the mutant nucleic acid sequence.

- 19. A mutant nucleic acld sequence produced by the method of claim 18.
- 20. A method for producing a polypeptide, comprising: (a) cultivating a strain comprising the mutant nucleic acid sequence of claim 19 encoding the polypeptide under conditions suitable for production of the polypeptide; and (b) recovering the polypeptide from the strain or supernatant, or recovering the strain containing the polypeptide.
- 21. A method for producing the polypeptide of claim 1 comprising: (a) cultivating a strain to produce a supernatant comprising the polypeptide; and (b) recovering the polypeptide from the strain or supernatant, or recovering the strain containing the polypeptide.
- A method for producing the polypeptide of claim 1 comprising: (a) cultivating a host cell comprising a nucleic acid construct comprising a nucleic acid sequence encoding the polypeptide under conditions suitable for production of the polypeptide; and (b) recovering the polypeptide from the host cell or supernatant, or recovering the host cell containing the polypeptide.